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=> file reg

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STRUCTURE FILE UPDATES: 26 SEP 2008 HIGHEST RN 1053621-88-7 DICTIONARY FILE UPDATES: 26 SEP 2008 HIGHEST RN 1053621-88-7

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=> s ?cyclovir

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If you are searching in a field that uses implied proximity, and you used a truncation symbol after a punctuation mark, the system may interpret the truncation symbol as being at the beginning of a term. Implied proximity is used in search fields indexed as single words, for example, the Basic Index.

=> file medicine

FILE 'DRUGMONOG' ACCESS NOT AUTHORIZED

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SINCE FILE TOTAL ENTRY SESSION 5.61 5.82

FULL ESTIMATED COST

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FILE 'USPATOLD' ENTERED AT 14:27:35 ON 28 SEP 2008
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=> s ll or ?cyclovir

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LEFT TRUNCATION IGNORED FOR FILE 'LIFESCI'
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 33 FILES SEARCHED...
        108103 L1 OR ?CYCLOVIR
Left truncation is not valid in the specified search field in the
specified file. The term has been searched without left truncation.
Examples: '?TERPEN?' would be searched as 'TERPEN?' and '?FLAVONOID'
would be searched as 'FLAVONOID.'
If you are searching in a field that uses implied proximity, and you
used a truncation symbol after a punctuation mark, the system may
interpret the truncation symbol as being at the beginning of a term.
Implied proximity is used in search fields indexed as single words,
for example, the Basic Index.
=> s psoriasis/ab
'AB' IS NOT A VALID FIELD CODE
L3
         99157 PSORIASIS/AB
=> s 12 and 13
T.4
          127 L2 AND L3
=> s valacyclovir
L5
         5458 VALACYCLOVIR
=> s 14 and 15
L6
             5 L4 AND L5
=> dup rem
ENTER L# LIST OR (END):16
DUPLICATE IS NOT AVAILABLE IN 'ADISINSIGHT, ADISNEWS, DGENE, DRUGMONOG2,
IMSPRODUCT, KOSMET, NUTRACEUT, PCTGEN, PHARMAML, USGENE'.
ANSWERS FROM THESE FILES WILL BE CONSIDERED UNIQUE
PROCESSING COMPLETED FOR L6
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              5 DUP REM L6 (0 DUPLICATES REMOVED)
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=> d 17 1-5 ibib, kwic

ANSWER 1 OF 5 CAPLUS COPYRIGHT 2008 ACS on STN

ACCESSION NUMBER: 2007:670138 CAPLUS <<LOGINID::20080928>>

DOCUMENT NUMBER: 147:102133

TITLE: Compositions and methods for treating dermatological

KIND DATE APPLICATION NO. DATE

conditions

Zhang, Jie; Warner, Kevin S.; Sharma, Sanjay INVENTOR(S):

PATENT ASSIGNEE(S): Zars, Inc., USA SOURCE: PCT Int. Appl., 74pp.

CODEN: PIXXD2

DOCUMENT TYPE: Patent LANGUAGE: English

FAMILY ACC. NUM. COUNT: 18

PATENT INFORMATION:

PATENT NO.

	2007				A2		2007 2008	0621		WO 2	006-	us47	747		2	0061	214
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		CF,	CG,	CI,	CM,	GΑ,	GN,	GQ,	GW,	ML,	MR,	ΝE,	SN,	TD,	ΤG,	BW,	GH,
		GM,	KE,	LS,	MW,	MZ,	NA,	SD,	SL,	SZ,	TZ,	UG,	ZM,	ZW,	AM,	AZ,	BY,
		KG,	KΖ,	MD,	RU,	ТJ,	TM,	AP,	EA,	EP,	OA						
AU	2006	3263	88		A1		2007	0621		AU 2	006-	3263	88		2	0061	214
AU	2006	3393	50		A1			0907								0061	214
EP	1959	929			A2		2008	0827		EP 2	006-	8476	57		2	0061	214
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significant penetration of the active. . . skin, which was greater than

the marketed Zovirax cream. The combination of isostearic acid and

trolamine enhanced the flux of acyclovir. The formulation

showed a sustained delivery of $\underline{acyclovir}$ for up to 8 h. It is reasonable to assume based on the drug load and the continued presence of the non-volatile solvent that the delivery of $\underline{acyclovir}$ would continue at the reported flux values for as long as the subject desires to leave the adhesive solidifying formulation. . .

50-03-3, Hydrocortisone acetate 50-23-7, Hydrocortisone 50-70-4, Sorbitol, biological studies 52-01-7, Spironolactone 54-42-2, Idoxuridine 56-81-5, Glycerol, biological studies 56-81-5D, Glycerol, fatty acid esters 57-13-6, Urea, biological studies 57-15-8, Chlorobutanol 57-55-6, Propylene glycol, biological studies 57-55-6D, Propylene glycol, fatty acid esters 60-29-7, Diethyl ether, biological studies 60-54-8, Tetracycline 64-17-5, Ethanol, biological studies 65-85-0, Benzoic acid, biological studies 67-56-1, Methanol, biological studies 67-63-0, Isopropanol, biological studies 67-64-1, Acetone, biological studies 67-68-5, Dimethyl sulfoxide, biological studies 67-73-2, Fluocinolone acetonide 67-97-0, Vitamin D3 68-12-2, Dimethylformamide, biological studies 69-65-8, Mannitol 70-00-8, Trifluridine 71-23-8, Propanol, biological studies 71-36-3, Butanol, biological studies 72-17-3, Sodium lactate 74-98-6, Propandiological studies 75-37-6, 1,1-Difluoroethane 76-25-5, Tracetonide 77-86-1, Tromethamine 77-93-0, Triethyl citrate 72-17-3, Sodium lactate 74-98-6, Propane, 75-37-6, 1,1-Difluoroethane 76-25-5, Triamcinolone Methyl ethyl ketone, biological studies 79-10-7D, Acrylic acid, derivs. 79-41-4, Methacrylic acid, biological studies 79-41-4D, Methacrylic acid, derivs., polymers 84-66-2, Diethyl phthalate 87-99-0, Xylitol 96-33-3, Methyl acrylate 97-00-7, Dinitrochlorobenzene 97-53-0, Eugenol 97-59-6, Allantoin 98-79-3, Pyroglutamic acid 100-51-6, Benzyl alcohol, biological studies 102-60-3, Neutrol TE 102-71-6, Trolamine, biological studies 102-76-1, Triacetin 104-46-1, p-Propenylanisole 104-55-2, Cinnamaldehyde 106-69-4, 1,2,6-Hexanetriol 106-97-8, Butane, biological studies 107-21-1, Ethylene glycol, biological studies 108-05-4, Vinyl acetate, biological studies 108-95-2, Phenol, biological studies 109-43-3, Dibutyl sebacate 109-66-0, Pentane, biological studies 110-16-7D, Maleic acid, copolymers 110-27-0, Isopropyl myristate 110-40-7, Diethyl sebacate 110-54-3, Hexane, biological studies 110-97-4, Diisopropanol amine 111-02-4. Squalene 111-42-2, Diethanolamine, biological studies 111-62-6, Ethyl oleate 111-90-0, Diethylene glycol monoethyl ether 112-38-9D, Undecylenic acid, derivs. 112-72-1, Myristyl alcohol 112-80-1, Oleic acid, biological studies 114-07-8, Erythromycin 115-10-6, Dimethyl ether 115-11-7, Isobutene, biological studies 120-40-1, Lauric diethanolamide 120-51-4, Benzyl benzoate 123-39-7, N-Methylformamide 123-92-2, Isoamyl acetate 126-07-8, Griseofulvin 131-11-3, Dimethyl phthalate 138-86-3, Limonene 141-78-6, Ethyl acetate, biological studies 142-91-6, Isopropyl palmitate 143-28-2, Oleyl alcohol 149-32-6, Erythrit 151-41-7, Lauryl sulfate 294-40-6, Cyclopentasiloxane 302-79-4, Tretinoin 356-12-7, Fluocinonide 382-67-2, Desoximethasone 431-89-0, 1,1,1,2,3,3,3-Heptafluoropropane 515-98-0, Ammonium lactate 518-28-5, Podofilox 585-86-4, Lactitol 585-88-6, Maltitol 638-94-8, Desonide 646-06-0D, Dioxolane, alkyl derivs. 661-19-8, Behenyl alcohol 676-46-0, Sodium malate 690-39-1, 1,1,1,3,3,3-Hexafluoropropane 768-94-5, Amantadine 777-11-7, Haloprogin 811-97-2, 1,1,1,2-Tetrafluoroethane 872-50-4, N-Methylpyrrolidone, biological studies 886-38-4, Diphenylcyclopropenone 996-31-6, Potassium lactate 1143-38-0, Anthralin 1314-13-2, Zinc oxide, biological studies 1320-51-0, Hydroxyethyl urea 1338-39-2, Sorbitan monolaurate 1338-43-8, Sorbitan monooleate 1397-89-3, Amphotericin B 1400-61-9, Nystatin 1404-04-2, Neomycin 1404-26-8, Polymyxin B 1405-87-4, Bacitracin 1406-18-4, Vitamin E 1524-88-5, Flurandrenolide 1984-15-2, Medronic acid 2002-29-1, Flumethasone pivalate 2152-44-5, Betamethasone valerate 2398-96-1, Tolnaftate

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2892-62-8, Squaric acid dibutyl ester 3056-17-5, Stavudine
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Halcinonide 4070-80-8 4759-48-2, Isotretinoin 5306-85-4, Dimethyl
isosorbide 5593-20-4, Betamethasone dipropionate 6283-92-7, Lauryl
lactate 7481-89-2, Zalcitabine 7681-93-8, Pimaricin 7732-18-5,
Water, biological studies 8011-96-9, Calamine 9000-07-1, Carrageenan 9000-30-0, Guar gum 9000-40-2, Locust bean gum 9002-88-4D,
Polyethylene, oxidized 9002-89-5, Polyvinyl alcohol 9003-01-4
9003-20-7, Polyvinyl acetate 9003-39-8, Polyvinylpyrrolidone
9003-70-7, Divinylbenzene-styrene copolymer 9004-32-4, Carboxymethyl
cellulose sodium 9004-34-6, Cellulose, biological studies 9004-35-7
9004-38-0, Cellulose acetate phthalate 9004-53-9, Dextrin 9004-57-3,
Ethyl cellulose 9004-62-0, Hydroxyethyl cellulose 9004-64-2,
Hydroxypropyl cellulose 9004-65-3, Hydroxypropyl methyl cellulose
9004-67-5, Methyl cellulose 9004-81-3, Polyethylene glycol laurate
9004-95-9 9004-96-0, Polyethylene glycol oleate 9004-99-3, PEG
stearate 9005-00-9 9005-07-6, Polyethylene glycol dioleate
9005-25-8, Starch, biological studies 9005-63-4, Polyoxyethylene
sorbitan 9005-63-4D, Polyethylene glycol sorbitan, fatty acid esters 9005-67-8 9006-65-9, Dimethicone 9011-16-9, Maleic anhydride-methyl vinyl ether copolymer 9011-16-9D, Maleic anhydride-vinyl methyl ether
copolymer, esters 9011-21-6, PEG glyceryl stearate 9063-89-2,
PEG-octyl phenyl ether 11138-66-2, Xanthan gum 12441-09-7D, Sorbitan,
fatty acid esters 12650-69-0, Mupirocin 13392-28-4, Rimantadine
13609-67-1, Hydrocortisone butyrate 14807-96-6, Talc, biological studies
16057-43-5, 2-[2-(Octadecyloxy)ethoxy]ethanol 16325-47-6 18323-44-9,
Clindamycin 22916-47-8, Miconazole 23593-75-1, Clotrimazole
24937-78-8, Ethylene-vinyl acetate copolymer 24938-16-7,
Dimethylaminoethyl methacrylate-butyl methacrylate-methyl methacrylate
copolymer 25013-16-5, Butylated hydroxyanisole 25087-26-7,
Poly(methacrylic acid) 25122-41-2, Clobetasol 25122-46-7, Clobetasol
propionate 25212-88-8, Methacrylic acid-ethyl acrylate copolymer
25265-71-8, Dipropylene glycol 25265-75-2, Butylene glycol 25322-68-3,
Polyethylene glycol 25322-68-3D, PEG, alkyl ethers and fatty acid esters
25322-69-4, Polypropylene glycol 25395-31-7, Diacetin 25496-72-4, Glycerol monooleate 25608-33-7, Butyl methacrylate-methyl methacrylate
copolymer 25618-55-7, Polyglycerol 26266-57-9, Sorbitan monopalmitate
26446-35-5, Monoacetin 27214-38-6, Glyceryl monomyristate 27220-47-9, Econazole 28874-51-3 30399-84-9, Isostearic acid 30516-87-1,
Zidovudine 31694-55-0D, triesters with fatty acids 32222-06-3,
Calcitriol 33434-24-1, Ethyl acrylate-methyl methacrylate-
{\tt trimethylammonioethyl\ methacrylate\ chloride\ copolymer}\qquad 33\,43\,4-2\,4-1
33564-31-7, Diflorasone diacetate 34513-50-3, Octyldodecanol
36653-82-4, Cetyl alcohol 36791-04-5, Ribavirin 37321-65-6, Propylene
glycol stearate 37353-59-6, Hydroxymethyl cellulose 38304-91-5,
Minoxidil 39809-25-1, Penciclovir 41621-49-2, Ciclopiroxolamine
51022-69-6, Amcinonide 53237-50-6 54182-62-6, Polacrilin
                                                                   54578-91-5,
Gantrez ES 425 56275-01-5 57107-95-6 57333-96-7, Tacalcitol
57524-89-7, Hydrocortisone valerate 59227-89-3, Azone 59277-89-3
, <u>Acyclovir</u> 59865-13-3, Cyclosporin 61318-90-9, Sulconazole
64211-45-6, Oxiconazole 64519-82-0, Isomalt 64872-76-0, Butoconazole 65277-42-1, Ketoconazole 65472-88-0, Naftifine 65899-73-2, Tioconazole
66734-13-2, Alclometasone dipropionate 66852-54-8, Halobetasol
propionate 67352-02-7 67915-31-5, Terconazole 68424-04-4,
Polydextrose 69655-05-6, Didanosine 75537-01-8, Gantrez S 97
RL: THU (Therapeutic use); BIOL (Biological study); USES (Uses)
   (solidifying adhesive compns. for treating dermatol. conditions)
78474-45-0, Plastoid B 78613-35-1, Amorolfine
                                                    80474-14-2, Fluticasone
propionate 82410-32-0, Gancyclovir 83919-23-7,
Mometasone furoate 84625-61-6, Itraconazole 85721-33-1, Ciprofloxacin
86386-73-4, Fluconazole 91161-71-6, Terbinafine 98319-26-7,
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Finasteride 99011-02-6, Imiquimod 101828-21-1, Butenafine
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    <u>Valacyclovir</u> 127779-20-8, Saquinavir 129618-40-2, Nevirapine 134678-17-4, Lamivudine 135668-52-9, Dermacryl 79 136470-78-5,
     Abacavir 136817-59-9, Delavirdine 137071-32-0, Picrolimus
     137234-62-9, Voriconazole 139110-80-8, Zanamivir 143780-36-3, Ethylene
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     Indinavir 154598-52-4, Efavirenz 155213-67-5, Ritonavir 158820-14-5
     159989-64-7, Nelfinavir 161814-49-9, Amprenavir 162808-62-0,
     Caspofungin 166663-25-8, Anidulafungin 171228-49-2, Posaconazole
     171664-79-2, Lactic acid-L-lactic acid copolymer 182760-06-1,
     Ravuconazole 195868-36-1, Phenyl trimethicone 196618-13-0, Oseltamivir
     227755-70-6 235114-32-6, Micafungin 357263-71-9, Honey Quat 50
     855659-57-3
     RL: THU (Therapeutic use); BIOL (Biological study); USES (Uses)
        (solidifying adhesive compns. for treating dermatol. conditions)
   ANSWER 2 OF 5 USPATFULL on STN
ACCESSION NUMBER:
                       2004:109827 USPATFULL <<LOGINID::20080928>>
                       Compositions and methods for treating inflammatory
TITLE:
                       diseases of the skin
                       Rothbard, Jonathan B., Cupertino, CA, United States
INVENTOR(S):
                       Wender, Paul A., Menlo Park, CA, United States
                       McGrane, P. Leo, Mountain View, CA, United States
                       Sista, Lalitha V. S., Sunnyvale, CA, United States
                       Kirschberg, Thorsten A., Mountain View, CA, United
                       States
PATENT ASSIGNEE(S):
                       CellGate, Inc., Sunnyvale, CA, United States (U.S.
                       corporation)
                          NUMBER
                                        KIND DATE
PATENT INFORMATION:
                       US 6730293 B1 20040504 US 2000-645689 20000824 (9)
APPLICATION INFO.:
                            NUMBER
                                          DATE
                       _____
PRIORITY INFORMATION: US 1999-150510P 19990824 (60)
DOCUMENT TYPE: Utility
FILE SEGMENT: GRANTED
PRIMARY EXAMINER: Webman, Edward J.
LEGAL REPRESENTATIVE: Townsend and Townsend and Crew LLP
NUMBER OF CLAIMS: 23
                      1
EXEMPLARY CLAIM:
NUMBER OF DRAWINGS: 31 Drawing Figure(s); 23 Drawing Page(s)
LINE COUNT:
                       2967
CAS INDEXING IS AVAILABLE FOR THIS PATENT.
       . . . as hydrocortisone, cyclosporin and FK506 across into and across
       one or more layers of the skin for the treatment of psoriasis
       and other inflammatory diseases of the skin.
      . . limited to, azole antifungals such as itraconazole, myconazole
DETD
      and fluconazole. Examples of antiviral agents include, but are not
      limited to, acyclovir, famciclovir, and valacyclovir
       . Such agents are useful for treating viral diseases, e.g., herpes.
      51-21-8, 5 Fluorouracil 58-08-2, Caffeine, biological studies
      60-54-8D, Tetracycline, derivs. 65-45-2, Salicylamide 69-53-4,
      Ampicillin 69-72-7, Salicylic acid, biological studies 100-33-4,
      Pentamidine 110-86-1D, Pyridine, trifluoro derivs., biological studies
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147-85-3, Proline, biological studies 1403-66-3, Gentamicin
     1406-05-9, Penicillin 1406-18-4, Vitamin e 9004-10-8, Insulin,
                                                    11111-12-9, Cephalosporin
     biological studies 11000-17-2, Vasopressin
     16110-51-3, Cromolyn 22916-47-8, Miconazole
                                                    57014-02-5, Eel
     calcitonin 58822-25-6, Leucine enkephalin 59277-89-3,
     Acyclovir 69049-73-6, Nedocromil 79217-60-0, Cyclosporin
      82410-32-0, Ganciclovir 84625-61-6, Itraconazole 86386-73-4,
     Fluconazole
        (compns. and methods for enhancing drug delivery across and into
        epithelial tissues)
    ANSWER 3 OF 5
                      MEDLINE on STN
ACCESSION NUMBER:
                   2004070493
                               MEDLINE <<LOGINID::20080928>>
DOCUMENT NUMBER:
                   PubMed ID: 14872168
TITLE:
                   Drug approval highlights for 2003.
AUTHOR:
                   Laustsen Gary; Wimett Lynn
CORPORATE SOURCE:
                   Regis University, Denver, Colorado, USA.
                   The Nurse practitioner, (2004 Feb) Vol. 29, No. 2, pp.
SOURCE:
                   8-15, 19-21; quiz 21-3. Ref: 28
                   Journal code: 7603663. ISSN: 0361-1817.
PUB. COUNTRY:
                   United States
                   Journal; Article; (JOURNAL ARTICLE)
DOCUMENT TYPE:
                   General Review; (REVIEW)
LANGUAGE:
                   English
                   Priority Journals; Nursing Journals
FILE SEGMENT:
ENTRY MONTH:
                   200404
                   Entered STN: 12 Feb 2004
ENTRY DATE:
                   Last Updated on STN: 1 May 2004
                   Entered Medline: 30 Apr 2004
     . . hormone therapy), Uroxatral (for benign prostatic hypertrophy),
AB
    Levitra (for erectile dysfunction), Flumist (for preventing influenza),
    Xolair (for asthma), Raptiva (for psoriasis), Cubicin (for skin
     infections), Crestor (for hypercholesterolemia), and Coreg (for severe
    heart failure).
    Check Tags: Female; Male
СТ
       *Acyclovir: AA, analogs & derivatives
        Acyclovir: TU, therapeutic use
     *Cardiovascular Agents: TU, therapeutic use
     Cardiovascular Diseases: DT, drug therapy
     *Dermatologic Agents: TU, therapeutic use
     *Drug.
    124832-27-5 (valacyclovir); 5633-20-5 (oxybutynin);
    59277-89-3 (Acyclovir); 7004-03-7 (Valine)
    ANSWER 4 OF 5 TOXCENTER COPYRIGHT 2008 ACS on STN
L7
ACCESSION NUMBER: 2004:114970 TOXCENTER <<LOGINID::20080928>>
COPYRIGHT:
                    Copyright (c) 2008 The Thomson Corporation
DOCUMENT NUMBER:
                    41-09588
TITLE:
                    Drug approval highlights for 2003
AUTHOR(S):
                    Laustsen, G; Wimett, L
SOURCE:
                    Nurse Practitioner (USA), (2004) Vol. 29, pp.
                    8-11,14-15,19-21. 28 Refs.
                    CODEN: NRPRDJ. ISSN: 0361-1817.
DOCUMENT TYPE:
                    Journal
FILE SEGMENT:
                    IPA
                    IPA 2004:9587
OTHER SOURCE:
LANGUAGE:
                    English
ENTRY DATE:
                    Entered STN: 25 May 2004
                    Last Updated on STN: 25 May 2004
AB. . . hormone therapy), Uroxatral (for benign prostatic hypertrophy),
```

```
Levitra (for erectile dysfunction), Flumist (for preventing influenza),
     Xolair (for asthma), Raptiva (for psoriasis), Cubicin (for skin
     infections), Crestor (for hypercholes-terolemia), and Coreg (for severe
     heart failure).
    Miscellaneous Descriptors
        Oxybutynin; urinary incontinence
         Valacyclovir; Herpes zoster
        Alfuzosin; prostatic hyperplasia
        Vardenafil; impotence
        Influenza vaccines; approvals
        Efalizumab; psoriasis
        Daptomycin; staphylococcal infections
        Carvedilol; heart failure
        Omalizumab; asthma
        Rosuvastatin calcium; hypercholesterolemia
        Estradiol diacetate; postmenopause
        Urinary incontinence; oxybutynin
        Parasympatholytic agents; oxybutynin
        Herpes zoster; valacyclovir
        Antivirals; valacyclovir
        Prostatic hyperplasia; alfuzosin
        Sympatholytic agents; alfuzosin
        Impotence; vardenafil
        Vasodilating agents; vardenafil
        Influenza; immunization
        Immunization; influenza
        Vaccines; influenza
        Immunomodulating agents; efalizumab
        Psoriasis; efalizumab
        Staphylococcal infections; daptomycin
        Antibiotics; daptomycin
        Heart failure; carvedilol
        Cardiac drugs; carvedilol
       Asthma;. .
    5633-20-5 (Oxybutynin)
RN
       124832-26-4 (Valacyclovir)
     81403-80-7 (Alfuzosin)
     224785-90-4 (Vardenafil)
     (Influenza vaccines)
     214745-43-4 (Efalizumab)
     103060-53-3 (Daptomycin)
     72956-09-3 (Carvedilol)
     242138-07-4 (Omalizumab)
     147098-20-2 (Rosuvastatin calcium)
     3434-88-6 (Estradiol diacetate)
    Oxybutynin (Oxytrol); Omalizumab (Xolair); Valacyclovir
     (Valtrex); Alfuzosin (Uroxatral); Vardenafil (Levitra); Rosuvastatin
     calcium (Crestor); Estradiol diacetate (Femring); Carvedilol (Coreg);
     Influenza vaccines (FluMist); Efalizumab (Raptiva); Daptomycin (Cubicin)
    ANSWER 5 OF 5 USPATFULL on STN
ACCESSION NUMBER:
                        2003:214302 USPATFULL <<LOGINID::20080928>>
TITLE:
                        Antimicrobial and anti-inflammatory peptides
                        McNicol, Patricia J., Vancouver, CANADA
INVENTOR(S):
                        Pawlak, Sonia K., Vancouver, CANADA
                        Rubinchik, Evelina, Richmond, CANADA
                        Cameron, Dale, Richmond, CANADA
                        Guarna, Maria Marta, Vancouver, CANADA
PATENT ASSIGNEE(S):
                       MICROLOGIX BIOTECH INC., Vancouver, CANADA, V6S 2L2
                        (non-U.S. corporation)
```

```
NUMBER
                                        KIND DATE
                       ______
PATENT INFORMATION: US 20030148945 A1 20030807 APPLICATION INFO.: US 2002-229368 A1 20020826
                                         A1 20020826 (10)
                              NUMBER
                                           DATE
                       _____
PRIORITY INFORMATION: US 2001-315003P 20010824 (60)
DOCUMENT TYPE:
                      Utility
FILE SEGMENT:
                      APPLICATION
LEGAL REPRESENTATIVE: SEED INTELLECTUAL PROPERTY LAW GROUP PLLC, 701 FIFTH
                     AVE, SUITE 6300, SEATTLE, WA, 98104-7092
                      25
NUMBER OF CLAIMS:
EXEMPLARY CLAIM:
                      1
                      2 Drawing Page(s)
NUMBER OF DRAWINGS:
LINE COUNT:
                       2380
CAS INDEXING IS AVAILABLE FOR THIS PATENT.
      . . . peptides are cationic peptides. The peptides are useful for the
       treatment of inflammatory diseases, such as microorganism-caused
       infections, acne, and \underline{\text{psoriasis.}} The peptides and peptide
       formulations may be used topically or parenterally.
      . . . also be used in combination with anti-viral agents. Exemplary
DETD
       anti-viral agents include, but are not limited to, amantadine
      hydrochloride, rimantadin, acyclovir, famciclovir, foscarnet,
       ganciclovir sodium, idoxuridine, ribavirin, sorivudine, trifluoridine,
       valacyclovir, vidarabin, didanosine, stavudine, zalcitabine,
       zidovudine, interferon alpha, and edoxudine.
=> s acyclovir
     63066 ACYCLOVIR
1.8
=> d his
     (FILE 'HOME' ENTERED AT 14:26:41 ON 28 SEP 2008)
     FILE 'REGISTRY' ENTERED AT 14:27:22 ON 28 SEP 2008
            46 S ?CYCLOVIR
T.1
     FILE 'ADISCTI, ADISINSIGHT, ADISNEWS, BIOSIS, BIOTECHNO, CAPLUS, DDFB,
     DGENE, DISSABS, DRUGB, DRUGMONOG2, DRUGU, EMBAL, EMBASE, ESBIOBASE,
     IFIPAT, IMSDRUGNEWS, IMSPRODUCT, IPA, KOSMET, LIFESCI, MEDLINE,
    NAPRALERT, NLDB, NUTRACEUT, PASCAL, PCTGEN, PHARMAML, ...' ENTERED AT
    14:27:35 ON 28 SEP 2008
        108103 S L1 OR ?CYCLOVIR
T.2
L3
         99157 S PSORIASIS/AB
L4
           127 S L2 AND L3
L5
          5458 S VALACYCLOVIR
L6
             5 S L4 AND L5
             5 DUP REM L6 (0 DUPLICATES REMOVED)
L7
1.8
         63066 S ACYCLOVIR
=> s 18 and 14
1.9
   74 L8 AND L4
=> s 19 and <20020809
MISSING TERM 'AND <20020809'
The search profile that was entered contains a logical
operator followed immediately by another operator.
```

```
=> s 19 and <2002
MISSING TERM 'AND <2002'
The search profile that was entered contains a logical
operator followed immediately by another operator.
=> s 19 <20020809
NUMERIC EXPRESSION NOT VALID 'L225 <20020809'
Numeric search expressions contain an operator (=,>,<,=<,=>), a field
qualifier, and the number or a range to be searched. Examples of
valid expressions are 'LD>6', '260-280/MW', and '10 < LD < 30'. For a
list of field codes in the current file, enter "HELP SFIELDS" at an
arrow prompt (=>). For more information on searching in numeric
fields, enter "HELP NUMERIC".
=> s 19 < 2002
NUMERIC EXPRESSION NOT VALID 'L225<2002'
Numeric search expressions contain an operator (=,>,<,=<,=>), a field
qualifier, and the number or a range to be searched. Examples of
valid expressions are 'LD>6', '260-280/MW', and '10 < LD < 30'. For a
list of field codes in the current file, enter "HELP SFIELDS" at an
arrow prompt (=>). For more information on searching in numeric
fields, enter "HELP NUMERIC".
=> s 19 < 2002
NUMERIC EXPRESSION NOT VALID 'L225 < 2002'
Numeric search expressions contain an operator (=,>,<,=<,=>), a field
qualifier, and the number or a range to be searched. Examples of
valid expressions are 'LD>6', '260-280/MW', and '10 < LD < 30'. For a
list of field codes in the current file, enter "HELP SFIELDS" at an
arrow prompt (=>). For more information on searching in numeric
fields, enter "HELP NUMERIC".
=> d his
     (FILE 'HOME' ENTERED AT 14:26:41 ON 28 SEP 2008)
     FILE 'REGISTRY' ENTERED AT 14:27:22 ON 28 SEP 2008
             46 S ?CYCLOVIR
T.1
     FILE 'ADISCTI, ADISINSIGHT, ADISNEWS, BIOSIS, BIOTECHNO, CAPLUS, DDFB,
     DGENE, DISSABS, DRUGB, DRUGMONOG2, DRUGU, EMBAL, EMBASE, ESBIOBASE,
     IFIPAT, IMSDRUGNEWS, IMSPRODUCT, IPA, KOSMET, LIFESCI, MEDLINE,
     NAPRALERT, NLDB, NUTRACEUT, PASCAL, PCTGEN, PHARMAML, ...' ENTERED AT
     14:27:35 ON 28 SEP 2008
         108103 S L1 OR ?CYCLOVIR
1.2
L3
          99157 S PSORIASIS/AB
L4
           127 S L2 AND L3
L5
           5458 S VALACYCLOVIR
              5 S L4 AND L5
1.6
L7
              5 DUP REM L6 (0 DUPLICATES REMOVED)
1.8
          63066 S ACYCLOVIR
             74 S L8 AND L4
L9
=> s 19 and pd<2002
   6 FILES SEARCHED...
'2002' NOT A VALID FIELD CODE
'2002' NOT A VALID FIELD CODE
'2002' NOT A VALID FIELD CODE
 15 FILES SEARCHED...
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'2002' NOT A VALID FIELD CODE

22 FILES SEARCHED...

'2002' NOT A VALID FIELD CODE

'2002' NOT A VALID FIELD CODE

'2002' NOT A VALID FIELD CODE

30 FILES SEARCHED...

32 FILES SEARCHED...

L10 28 L9 AND PD<2002

=> dup rem

ENTER L# LIST OR (END):110

DUPLICATE IS NOT AVAILABLE IN 'ADISINSIGHT, ADISNEWS, DGENE, DRUGMONOG2,

IMSPRODUCT, KOSMET, NUTRACEUT, PCTGEN, PHARMAML, USGENE'.

ANSWERS FROM THESE FILES WILL BE CONSIDERED UNIQUE

PROCESSING COMPLETED FOR L10

L11 22 DUP REM L10 (6 DUPLICATES REMOVED)

 \Rightarrow d 111 1-22 ibib, kwic

L11 ANSWER 1 OF 22 CAPLUS COPYRIGHT 2008 ACS on STN DUPLICATE 1

ACCESSION NUMBER: 2001:719000 CAPLUS <<LOGINID::20080928>>

DOCUMENT NUMBER: 135:262277

TITLE: Pharmaceutical compositions and methods for managing

skin conditions

INVENTOR(S):
Murad, Howard

PATENT ASSIGNEE(S): USA

SOURCE: U.S., 18 pp., Cont.-in-part of U.S. 6,071,541.

CODEN: USXXAM

DOCUMENT TYPE: Patent LANGUAGE: English

FAMILY ACC. NUM. COUNT: 4

PATENT INFORMATION:

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
US 6296880	B1	20011002	US 2000-549202	20000413 <
US 6071541	A	20000606	US 1999-330127	19990611 <
US 20020041901	A1	20020411	US 2001-878231	20010612
US 6383523	В2	20020507		
US 20030007939	A1	20030109	US 2002-77928	20020220
US 20020172719	A1	20021121	US 2002-93443	20020311
US 7018660	В2	20060328		
US 20040091548	A1	20040513	US 2003-702453	20031107
US 20060051429	A1	20060309	US 2005-249496	20051014
PRIORITY APPLN. INFO.:			US 1998-94775P	P 19980731
			US 1999-330127	A2 19990611
			US 2000-549202	A1 20000413
			US 2001-878231	A2 20010612
			US 2001-953431	A2 20010917
			US 2002-93443	A1 20020311
REFERENCE COUNT:	29	THERE ARE 29		AILABLE FOR THIS IN THE RE FORMAT

PΙ	US 6296880 B1 200	11002					
	PATENT NO.	KIND	DATE	APPLICATION NO.	DATE		
ΡI	US 6296880	В1	20011002	US 2000-549202	20000413 <		
	US 6071541	A	20000606	US 1999-330127	19990611 <		
	US 20020041901	A1	20020411	US 2001-878231	20010612		
	US 6383523	В2	20020507				
	US 20030007939	A1	20030109	US 2002-77928	20020220		

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      US
      20020172719
      A1
      20021121
      US
      2002-93443
      20020311

      US
      7018660
      B2
      20060328
      US
      2003-702453
      20031107

      US
      20060051429
      A1
      20060309
      US
      2005-249496
      20051014

     . . . methods for the cleansing of skin to facilitate the prevention,
     treatment, and management of skin conditions, such as seborrheic
     dermatitis, psoriasis, folliculitis, rosacea, perioral
     dermatitis, acne, impetigo and other inflammatory skin conditions, and the
     like, including a sufficient amount of an. . .
     50-21-5, Lactic acid, biological studies 68-26-8, Retinol 69-72-7,
TΤ
     Salicylic acid, biological studies 77-92-9, Citric acid, biological
     studies 79-14-1, Glycolic acid, biological studies 557-34-6, Zinc
     acetate 1314-13-2, Zinc oxide, biological studies 3380-34-5, Triclosan
     7704-34-9, Sulfur, biological studies 7722-84-1, Hydrogen peroxide,
     biological studies 23593-75-1, Clotrimazole 39809-25-1, Penciclovir
     41621-49-2, Ciclopirox olamine 59277-89-3, Acyclovir
     68797-35-3, Dipotassium glycyrrhizate
     RL: THU (Therapeutic use); BIOL (Biological study); USES (Uses)
        (topical compns. for managing skin conditions containing acids and hydrogen
        peroxide and antivirals and other actives)
L11 ANSWER 2 OF 22 USPATFULL on STN
ACCESSION NUMBER:
                      2001:100886 USPATFULL <<LOGINID::20080928>>
TITLE:
                        Anti-inflammatory formulations for inflammatory
                         diseases
INVENTOR(S):
                        Kross, Robert D., Bellmore, NY, United States
                         Siff, Elliott J., Westport, CT, United States
PATENT ASSIGNEE(S):
                        Alcide Corporation, Redmond, WA, United States (U.S.
                         corporation)
                            NUMBER KIND DATE
                         -----
                         US 37263 E1 20010703
US 5384134 19950124
PATENT INFORMATION:
                                            19950124 (Original)
                         US 1997-787144 19970122 (8)
US 1993-115461 19930901 (Original)
APPLICATION INFO.:
                         Continuation of Ser. No. US 1992-930088, filed on 14
RELATED APPLN. INFO.:
                         Aug 1992, now abandoned Division of Ser. No. US
                         1990-543655, filed on 26 Jun 1990, now abandoned
                         Division of Ser. No. US 1988-202758, filed on 3 Jun
                         1988, now patented, Pat. No. US 4956184
                         Continuation-in-part of Ser. No. US 1988-190798, filed
                         on 6 May 1988, now abandoned
DOCUMENT TYPE:
                        Reissue
FILE SEGMENT:
FILE SEGMENT: GRANTED
PRIMARY EXAMINER: Criares, Theodore J.
LEGAL REPRESENTATIVE: Seed and Berry LLP
NUMBER OF CLAIMS: 10
                       1
4 Drawing Figure(s); 4 Drawing Page(s)
767
EXEMPLARY CLAIM:
NUMBER OF DRAWINGS:
LINE COUNT:
CAS INDEXING IS AVAILABLE FOR THIS PATENT.
AΒ
       There is disclosed a method for treating dermatologic diseases caused by
       microbial overgrowth or inflammation, such as psoriasis,
       fungal infections, eczema, dandruff, acne, genital herpes lesions, and
       leg ulcers. There is further disclosed an antiviral lubricating
       composition that. . .
       . . . and secondary attacks become less frequent with time.
SUMM
```

```
discomfort of the lesion. Acyclovir, applied topically, tends
      to decrease pain of the primary lesions, but it has not proven very
      effective for decreasing vital shedding or lesion duration. Topical
      acyclovir has not been shown to be particularly effective for
      reducing or treating recurrent disease.
SUMM
      Acyclovir is a purine nucleoside analog that is selectively
      cidal to the herpes simplex virus because only the thymidine kinase
      enzyme of herpes simplex virus can convert acyclovir to its
      monophosphate form while host cell thymidine kinase cannot. The
      monophosphate form is converted to an acyclovir triphosphate,
      which can interfere with vital DNA replication. Topical
      acyclovir is applied as a 5% ointment every three hours, or up
      to eight times daily, for at least seven days.. . . patient
      compliance problems for dosing in the genital areas throughout the day
      and throughout the night. A further problem of acyclovir has
      been the development resistant strains of herpes simplex, caused by a
      mutation of the thymidine kinase gene. Accordingly, no backup treatments
      are available for acyclovir-resistant herpes simplex
      infections. This problem exists with most antibiotic microbial
      treatments, but is generally not a problem non-antibiotic treatments.
      . . . as re-epithelialization of the original lesions). The results
DETD
      of the study were compared to a similar study conducted with topical
      acyclovir and placebo (Fiddian et al, J. Antimicrob. Chem.
      12:Suppl. B:67-77, 1983) and are presented together in Table 1 below:
           Symptoms Shedding Healing Rate
DETD
                                                  Median Recurrence
                        Time (d) Time (d)
                  1 * *
Example 1
           3 *
                            8 (1-17)
                                       19.4
(32)
                             7-8
 Acyclovir
                          3
                               10-13
Placebo
                        6-9
                                           55
*Twenty-one of twenty-four patients had a duration of symptoms of 5 or less. .
      twice daily dosing (compared with 5 times daily with some treatments
      such as Acyclovir)
L11 ANSWER 3 OF 22 CAPLUS COPYRIGHT 2008 ACS on STN DUPLICATE 2
ACCESSION NUMBER: 1997:527765 CAPLUS <<LOGINID::20080928>>
DOCUMENT NUMBER:
                       127:185859
ORIGINAL REFERENCE NO.: 127:35885a,35888a
TITLE:
                       Nucleotides for topical treatment of psoriasis
INVENTOR(S):
                       Hostetler, Karl Y.
PATENT ASSIGNEE(S):
                       USA
SOURCE:
                       U.S., 15 pp., Cont.-in-part of U.S. 5,580,571.
                       CODEN: USXXAM
DOCUMENT TYPE:
                       Patent
LANGUAGE:
                       English
FAMILY ACC. NUM. COUNT: 5
PATENT INFORMATION:
    PATENT NO.
                       KIND DATE APPLICATION NO.
                              -----
                                         _____
    _____
                       ____
    US 5654286
                       A
                             19970805 US 1995-485025 19950607 <--
    US 5580571
                       A
                             19961203 US 1993-60258
                                                              19930512 <--
                       A1 19961219
                                       CA 1996-222224
    CA 2222224
    WO 9640166
                       A1 19961219
                                       WO 1996-US10084
                                                              19960606 <--
        W: AL, AM, AT, AU, AZ, BB, BG, BR, BY, CA, CH, CN, CZ, DE, DK, EE,
            ES, FI, GB, GE, HU, IL, IS, JP, KE, KG, KP, KR, KZ, LK, LR, LS,
```

LT, LU, LV, MD, MG, MK, MN, MW, MX, NO, NZ, PL, PT, RO, RU, SD,

Treatments include drying agents to symptomatically lessen the

```
SE, SG
        RW: KE, LS, MW, SD, SZ, UG, AT, BE, CH, DE, DK, ES, FI, FR, GB, GR,
            IE, IT, LU, MC, NL, PT, SE, BF, BJ, CF, CG, CI, CM, GA, GN
    AU 9662737 A 19961230 AU 1996-62737 19960606 <---
EP 831855 A1 19980401 EP 1996-921531 19960606 <---
        R: AT, BE, CH, DE, DK, ES, FR, GB, GR, IT, LI, LU, NL, SE, MC, PT,
            IE, FI
     CN 1220605
                              19990623
                                          CN 1996-195944
                                                                19960606 <--
     JP 2002515018 T 20020521
                                          JP 1997-502193
                                                                 19960606
                                                            A2 19930512
PRIORITY APPLN. INFO.:
                                          US 1993-60258
                                                            B2 19911015
                                          US 1991-777683
                                           US 1995-485025
                                                            A 19950607
                                           WO 1996-US10084
                                                             W 19960606
OTHER SOURCE(S):
                       MARPAT 127:185859
    US 5654286 A <u>19970805</u>
    PATENT NO. KIND DATE APPLICATION NO.
                                                                DATE
                                        _____
                                                                -----
     _____
                        ----
    US 5654286 A 19970805 US 1995-485025 19950607 <---
US 5580571 A 19961203 US 1993-60258 19930512 <---
PΙ
                 A1 19961219 CA 1996-2222224 19960606 <--
A1 19961219 WO 1996-US10084 19960606 <--
     CA 2222224
     WO 9640166
        W: AL, AM, AT, AU, AZ, BB, BG, BR, BY, CA, CH, CN, CZ, DE, DK, EE,
            ES, FI, GB, GE, HU, IL, IS, JP, KE, KG, KP, KR, KZ, LK, LR, LS,
            LT, LU, LV, MD, MG, MK, MN, MW, MX, NO, NZ, PL, PT, RO, RU, SD,
            SE, SG
        RW: KE, LS, MW, SD, SZ, UG, AT, BE, CH, DE, DK, ES, FI, FR, GB, GR,
            IE, IT, LU, MC, NL, PT, SE, BF, BJ, CF, CG, CI, CM, GA, GN
    AU 9662737
                        A 19961230 AU 1996-62737
A1 19980401 EP 1996-921531
                                                                19960606 <--
     EP 831855
                                                                19960606 <--
        R: AT, BE, CH, DE, DK, ES, FR, GB, GR, IT, LI, LU, NL, SE, MC, PT,
            IE, FI
     CN 1220605
                   А
т
                                                                19960606 <--
                             19990623 CN 1996-195944
                        T 20020521 JP 1997-502193
     JP 2002515018
                                                                19960606
     Psoriasis and other diseases of skin cell hyperproliferation are
AΒ
     treated with topical pharmaceutical compns. containing mono-, di-, and
     triphosphate esters of. . .
     134-46-3, 5-Fluorodeoxyuridine monophosphate 796-66-7, 5-Fluorouridine
ΙT
     monophosphate 803-98-5, 5-Fluorouridine diphosphate 1049-56-5
     2018-19-1, 6-Azauridine monophosphate 2710-64-7, 5-Fluorodeoxyuridine
     triphosphate 3828-96-4, 5-Fluorouridine triphosphate 6198-30-7
     66004-77-1, 2',3'-Dideoxycytidine 5'-triphosphate 66341-18-2,
     Acyclovir triphosphate 70711-50-1 104086-75-1 104086-76-2
     104959 - 32 - 2 106867 - 30 - 5 134646 - 41 - 6 134646 - 42 - 7 167620 - 89 - 5
     180297-84-1
                 186553-15-1
     RL: BAC (Biological activity or effector, except adverse); BSU (Biological
     study, unclassified); THU (Therapeutic use); BIOL (Biological study); USES
     (Uses)
        (nucleotides for topical treatment of psoriasis)
L11 ANSWER 4 OF 22 USPATFULL on STN
ACCESSION NUMBER:
                      97:96851 USPATFULL <<LOGINID::20080928>>
                       Method of treating lesions resulting from genital
TITLE:
                       herpes with hyaluronic acid-urea pharmaceutical
                       compositions
INVENTOR(S):
                       Gallina, Damian J., Erie, PA, United States
                      Patent Biopharmaceutics, Inc., Erie, PA, United States
PATENT ASSIGNEE(S):
                       (U.S. corporation)
                          NUMBER KIND DATE
```

PATENT INFORMATION: US 5679655 19971021 APPLICATION INFO.: US 1995-471323 19950602 (8)

RELATED APPLN. INFO.: Division of Ser. No. US 1993-101826, filed on 4 Aug

1993

DOCUMENT TYPE: Utility FILE SEGMENT: Granted

PRIMARY EXAMINER: Lilling, Herbert J. LEGAL REPRESENTATIVE: Cushman Darby & Cushman

8 NUMBER OF CLAIMS: EXEMPLARY CLAIM: 1

NUMBER OF DRAWINGS: 6 Drawing Figure(s); 6 Drawing Page(s) 1447

LINE COUNT:

CAS INDEXING IS AVAILABLE FOR THIS PATENT.

AΒ . . . erythema, edema, papules, vesicles, macules, pustules, scaling, cracking, crusting, and lesions. The invention further includes methods for the treatment of psoriasis, eczema, dermatitis, herpetic conditions, acne, skin ulcers, genital herpes lesions and anorectal disease, which includes applying to tissues in need. . .

. . . routinely in a dermatologic office setting. Normally, these DETD conditions have been treated with various topical medications including topical corticosteroids, topical $\underline{acyclovir}$ and sometimes internal medications such as oral $\overline{corticosteroids}$. Patients also tend to treat skin lesions with many over-the-counter medications such. . .

L11 ANSWER 5 OF 22 USPATFULL on STN

97:42867 USPATFULL <<LOGINID::20080928>> ACCESSION NUMBER:

TITLE: Hyaluronic acid-urea pharmaceutical compositions

utilized for treatment of diseases of cutis

INVENTOR(S): Gallina, Damian J., Erie, PA, United States

PATENT ASSIGNEE(S): Patent Biopharmaceutics, Inc., Erie, PA, United States

(U.S. corporation)

NUMBER KIND DATE

US 5631242 19970520 US 1995-471334 19950602 (8) PATENT INFORMATION:

APPLICATION INFO.:

RELATED APPLN. INFO.: Division of Ser. No. US 1993-101826, filed on 4 Aug

1993

DOCUMENT TYPE: Utility FILE SEGMENT: Granted
PRIMARY EXAMINER: Lilling, Herbert J.

LEGAL REPRESENTATIVE: Cushman Darby & Cushman

NUMBER OF CLAIMS: 8 EXEMPLARY CLAIM: 1

NUMBER OF DRAWINGS: 6 Drawing Figure(s); 6 Drawing Page(s) LINE COUNT: 1451

CAS INDEXING IS AVAILABLE FOR THIS PATENT.

. . . erythema, edema, papules, vesicles, macules, pustules, scaling, cracking, crusting, and lesions. The invention further includes methods for the treatment of psoriasis, eczema, dermatitis, herpetic conditions, acne, skin ulcers, genital herpes lesions and anorectal disease, which includes applying to tissues in need. . .

DETD . . . routinely in a dermatologic office setting. Normally, these conditions have been treated with various topical medications including topical corticosteroids, topical acyclovir and sometimes internal medications such as oral corticosteroids. Patients also tend to treat skin lesions with many over-the-counter medications such. . .

L11 ANSWER 6 OF 22 USPATFULL on STN

ACCESSION NUMBER: 97:36175 USPATFULL <<LOGINID::20080928>>

TITLE: Hyaluronic acid-urea pharmaceutical compositions and

uses

INVENTOR(S): Gallina, Damian J., Erie, PA, United States

PATENT ASSIGNEE(S): Patent Biopharmaceutics, Inc., Erie, PA, United States

(U.S. corporation)

NUMBER KIND DATE ______

PATENT INFORMATION: US 5624915 19970429 APPLICATION INFO.: US 1995-471327 19950602 (8)

RELATED APPLN. INFO.: Division of Ser. No. US 1993-101826, filed on 4 Aug

1993

DOCUMENT TYPE: Utility

FILE SEGMENT: Granted PRIMARY EXAMINER: Lilling, Herbert J.

LEGAL REPRESENTATIVE: Cushman Darby & Cushman IP Group of Pillsbury Madison &

Sutro LLP

NUMBER OF CLAIMS: EXEMPLARY CLAIM:

6 Drawing Figure(s); 6 Drawing Page(s) 1450 NUMBER OF DRAWINGS:

LINE COUNT:

CAS INDEXING IS AVAILABLE FOR THIS PATENT.

. . . erythema, edema, papules, vesicles, macules, pustules, scaling, AB cracking, crusting, and lesions. The invention further includes methods

for the treatment of psoriasis, eczema, dermatitis, herpetic

conditions, acne, skin ulcers, genital herpes lesions and anorectal

disease, which includes applying to tissues in need. . .

DETD . . . routinely in a dermatologic office setting. Normally, these

conditions have been treated with various topical medications including

topical corticosteroids, topical $\underline{\mathtt{acyclovir}}$ and sometimes

internal medications such as oral corticosteroids. Patients also tend to treat skin lesions with many over-the-counter medications such. . .

L11 ANSWER 7 OF 22 CAPLUS COPYRIGHT 2008 ACS on STN DUPLICATE 3

ACCESSION NUMBER: 1997:132780 CAPLUS <<LOGINID::20080928>>

DOCUMENT NUMBER: 126:139875

ORIGINAL REFERENCE NO.: 126:26883a,26886a

TITLE: Nucleotide analogs, their preparation, and

pharmaceutical compositions containing them for

topical treatment of proliferative disease of the skin

INVENTOR(S): Hostetler, Karl Y.

PATENT ASSIGNEE(S): Hostetler, Karl Y., USA SOURCE: PCT Int. Appl., 31 pp.

CODEN: PIXXD2

DOCUMENT TYPE: Patent LANGUAGE: English

FAMILY ACC. NUM. COUNT: 5

PATENT INFORMATION:

KIND DATE APPLICATION NO. DATE PATENT NO. ----_____ ____ ______ WO 9640166 A1 19961219 WO 1996-US10084 19960606 <--W: AL, AM, AT, AU, AZ, BB, BG, BR, BY, CA, CH, CN, CZ, DE, DK, EE, ES, FI, GB, GE, HU, IL, IS, JP, KE, KG, KP, KR, KZ, LK, LR, LS, LT, LU, LV, MD, MG, MK, MN, MW, MX, NO, NZ, PL, PT, RO, RU, SD, SE, SG RW: KE, LS, MW, SD, SZ, UG, AT, BE, CH, DE, DK, ES, FI, FR, GB, GR,

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IE, IT, LU, MC, NL, PT, SE, BF, BJ, CF, CG, CI, CM, GA, GN
     US 5654286 A 19970805 US 1995-485025 19950607 <--
                         A 19961230 AU 1996-62737
A1 19980401 EP 1996-921531
                                                                  19960606 <--
                        Α
     AU 9662737
     EP 831855
                                                                   19960606 <--
         R: AT, BE, CH, DE, DK, ES, FR, GB, GR, IT, LI, LU, NL, SE, MC, PT,
             IE, FI
                              20020521
                                            JP 1997-502193
     JP 2002515018
                                                                   19960606
                                            US 1995-485025 A 19950607
US 1993-60258 A2 19930512
WO 1996-US10084 W 19960606
PRIORITY APPLN. INFO.:
OTHER SOURCE(S):
                       MARPAT 126:139875
    WO 9640166 A1 19961219
    PATENT NO.
                        KIND DATE APPLICATION NO.
                                          _____
     _____
                        ----
    WO 9640166
                        A1 19961219 WO 1996-US10084 19960606 <--
РΤ
        W: AL, AM, AT, AU, AZ, BB, BG, BR, BY, CA, CH, CN, CZ, DE, DK, EE,
             ES, FI, GB, GE, HU, IL, IS, JP, KE, KG, KP, KR, KZ, LK, LR, LS,
             LT, LU, LV, MD, MG, MK, MN, MW, MX, NO, NZ, PL, PT, RO, RU, SD,
             SE, SG
         RW: KE, LS, MW, SD, SZ, UG, AT, BE, CH, DE, DK, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE, BF, BJ, CF, CG, CI, CM, GA, GN
                         A 19970805 US 1995-485025 19950607 <---
A 19961230 AU 1996-62737 19960606 <---
     US 5654286
     AU 9662737
                        A1 19980401 EP 1996–921531 19960606 <--
     EP 831855
        R: AT, BE, CH, DE, DK, ES, FR, GB, GR, IT, LI, LU, NL, SE, MC, PT,
            IE, FI
     JP 2002515018
                              20020521
                                          JP 1997-502193
     . . . nucleoside analogs, DNA chain-terminating dideoxynucleoside
     analogs and other nucleoside analogs for the topical treatment of
     hyperproliferative diseases of the skin (\underline{psoriasis}, atopic
     dermatitis, basal cell carcinoma, etc.). The useful phosphate esters of
     the nucleoside analogs include phosphoramidates and phosphothiorates, as
     well. . .
     50-91-9DP, 5-Fluorodeoxyuridine, derivs. 54-25-1DP, 6-Azauridine,
ТТ
     derivs. 134-46-3P, 5-Fluorodeoxyuridine monophosphate 147-94-4DP,
     Cytosine arabinoside, derivs. 316-46-1DP, 5-Fluorouridine, derivs.
     342-69-8DP, 6-Methylmercaptopurine riboside, derivs. 796-66-7P,
     5-Fluorouridine monophosphate 803-98-5P, 5-Fluorouridine diphosphate
     1049-56-5P 2018-19-1P, 6-Azauridine monophosphate 2710-64-7P,
     5-Fluorodeoxyuridine triphosphate 3416-05-5DP, derivs. 3828-96-4P,
     5-Fluorouridine triphosphate 4097-22-7DP, Dideoxyadenosine, derivs.
     4291-63-8DP, 2-Chlorodeoxyadenosine, derivs. 6198-30-7P 7481-89-2DP,
     Dideoxycytidine, derivs. 20227-41-2DP, derivs. 38819-10-2DP, derivs.
     40627-14-3DP, derivs. 59277-89-3DP, Acyclovir, derivs.
     60129-59-1DP, 2'-Deoxytubercidin, derivs. 66004-77-1P
     <u>66341-18-2P</u>, <u>Acyclovir</u> triphosphate 69655-05-6DP,
     Dideoxyinosine, derivs. 70711-50-1P 82410-32-0DP, Ganciclovir,
     derivs. 85326-06-3DP, derivs. 104086-75-1P 104086-76-2P
     104904-94-1P 104904-96-3P 104959-32-2P, 2-Chloro-2'-deoxyadenosine monophosphate 106867-30-5P 123318-82-1DP, derivs. 134646-41-6P
    134646-42-7P 167620-89-5P 180297-84-1P 186553-12-8P 186553-15-1P
     186553-16-2P 186553-17-3P
    RL: BAC (Biological activity or effector, except adverse); BSU (Biological
     study, unclassified); SPN (Synthetic preparation); THU (Therapeutic use);
     BIOL (Biological study); PREP (Preparation); USES (Uses)
        (nucleotide analogs, preparation, and pharmaceutical compns. for topical
        treatment of proliferative skin diseases)
L11 ANSWER 8 OF 22 USPATFULL on STN
```

96:113915 USPATFULL <<LOGINID::20080928>>

ACCESSION NUMBER:

TITLE: Hyaluronic acid-urea pharmaceutical compositions and

uses

Gallina, Damian J., Erie, PA, United States INVENTOR(S):

PATENT ASSIGNEE(S): Patent Biopharmaceutics, Inc., Erie, PA, United States

(U.S. corporation)

NUMBER KIND DATE _____

US 5583120 US 5583120 19961210 US 1995-471332 19950602 (8) PATENT INFORMATION:

APPLICATION INFO.:

RELATED APPLN. INFO.: Division of Ser. No. US 1993-101826, filed on 4 Aug

1993

DOCUMENT TYPE: Utility FILE SEGMENT: Granted PRIMARY EXAMINER: Lilling, Herbert J.

LEGAL REPRESENTATIVE: Cushman Darby & Cushman IP Group of Pillsbury Madison &

Sutro LLP

NUMBER OF CLAIMS: EXEMPLARY CLAIM: 1

6 Drawing Figure(s); 6 Drawing Page(s) 1449 NUMBER OF DRAWINGS:

LINE COUNT:

CAS INDEXING IS AVAILABLE FOR THIS PATENT.

. . . erythema, edema, papules, vesicles, macules, pustules, scaling, AB cracking, crusting, and lesions. The invention further includes methods

for the treatment of psoriasis, eczema, dermatitis, herpetic

conditions, acne, skin ulcers, genital herpes lesions and anorectal

disease, which includes applying to tissues in need. . .

DETD . . . routinely in a dermatologic office setting. Normally, these conditions have been treated with various topical medications including

topical corticosteroids, topical acyclovir and sometimes

internal medications such as oral corticosteroids. Patients also tend to treat skin lesions with many over-the-counter medications such. . .

L11 ANSWER 9 OF 22 USPATFULL on STN

ACCESSION NUMBER: 96:113914 USPATFULL <<LOGINID::20080928>>

TITLE: Hyaluronic acid-urea pharmaceutical compositions and

uses

INVENTOR(S): Gallina, Damian J., Erie, PA, United States

Patent Biopharmaceutics, Inc., Erie, PA, United States PATENT ASSIGNEE(S):

(U.S. corporation)

NUMBER KIND DATE ______

PATENT INFORMATION: US 5583119 19961210 APPLICATION INFO.: US 1995-471330 19950602 (8)

RELATED APPLN. INFO.: Division of Ser. No. US 1993-101826, filed on 4 Aug

1993

DOCUMENT TYPE: Utility FILE SEGMENT: Granted

PRIMARY EXAMINER: Lilling, Herbert J.

LEGAL REPRESENTATIVE: Cushman Darby & Cushman IP Group of Pillsbury Madison &

Sutro LLP

NUMBER OF CLAIMS: EXEMPLARY CLAIM: 1

NUMBER OF DRAWINGS: 6 Drawing Figure(s); 6 Drawing Page(s)

LINE COUNT:

CAS INDEXING IS AVAILABLE FOR THIS PATENT.

. . erythema, edema, papules, vesicles, macules, pustules, scaling,

cracking, crusting, and lesions. The invention further includes methods for the treatment of psoriasis, eczema, dermatitis, herpetic conditions, acne, skin ulcers, genital herpes lesions and anorectal

disease, which includes applying to tissues in need. . .

DETD . . . routinely in a dermatologic office setting. Normally, these conditions have been treated with various topical medications including topical corticosteroids, topical <u>acyclovir</u> and sometimes internal medications such as oral corticosteroids. Patients also tend to treat skin lesions with many over-the-counter medications such. .

L11 ANSWER 10 OF 22 USPATFULL on STN

ACCESSION NUMBER: 96:113913 USPATFULL <<LOGINID::20080928>> Method of treating an anorectal disease using TITLE: hyaluronic acid-urea pharmaceutical compositions

Gallina, Damian J., Erie, PA, United States INVENTOR(S):

PATENT ASSIGNEE(S): Patent Biopharmaceutics, Inc., Erie, PA, United States

(U.S. corporation)

NUMBER KIND DATE ______

PATENT INFORMATION: US 5583118 19961210 APPLICATION INFO:: US 1995-458303 19950602 (8)

RELATED APPLN. INFO.: Division of Ser. No. US 1993-101826, filed on 4 Aug

1993

DOCUMENT TYPE: Utility FILE SEGMENT: Granted
PRIMARY EXAMINER: Lilling, Herbert J.

LEGAL REPRESENTATIVE: Cushman Darby & Cushman

NUMBER OF CLAIMS: 8 EXEMPLARY CLAIM: 1

NUMBER OF DRAWINGS: 6 Drawing Figure(s); 6 Drawing Page(s)

LINE COUNT: 1451

CAS INDEXING IS AVAILABLE FOR THIS PATENT.

. . . erythema, edema, papules, vesicles, macules, pustules, scaling, cracking, crusting, and lesions. The invention further includes methods for the treatment of psoriasis, eczema, dermatitis, herpetic

conditions, acne, skin ulcers, genital herpes lesions and anorectal

disease, which includes applying to tissues in need. . .

. . . routinely in a dermatologic office setting. Normally, these DETD conditions have been treated with various topical medications including topical corticosteroids, topical acyclovir and sometimes internal medications such as oral corticosteroids. Patients also tend to treat skin lesions with many over-the-counter medications such. . .

L11 ANSWER 11 OF 22 USPATFULL on STN

ACCESSION NUMBER: 96:77765 USPATFULL <<LOGINID::20080928>>

TITLE: Hyaluronic acid-urea pharmaceutical compositions and

uses

INVENTOR(S): Gallina, Damian J., Erie, PA, United States

Patent Biopharmaceutics, Inc., Erie, PA, United States PATENT ASSIGNEE(S):

(U.S. corporation)

NUMBER KIND DATE -----

PATENT INFORMATION: US 5550112 19960827 APPLICATION INFO:: US 1993-101826 19930804 (8)

RELATED APPLN. INFO.: Continuation-in-part of Ser. No. US 1992-966938, filed

on 30 Dec 1992, now abandoned

DOCUMENT TYPE: Utility

Granted FILE SEGMENT:

PRIMARY EXAMINER: Lilling, Herbert J. LEGAL REPRESENTATIVE: Cushman Darby & Cushman

NUMBER OF CLAIMS: EXEMPLARY CLAIM:

6 Drawing Figure(s); 6 Drawing Page(s) NUMBER OF DRAWINGS:

LINE COUNT: 1427

CAS INDEXING IS AVAILABLE FOR THIS PATENT.

. . . erythema, edema, papules, vesicles, macules, pustules, scaling, AB cracking, crusting, and lesions. The invention further includes methods for the treatment of psoriasis, eczema, dermatitis, herpetic conditions and acne, which includes applying to tissues in need of such treatment a therapeutically effective amount. . .

DETD . . . routinely in a dermatologic office setting. Normally, these conditions have been treated with various topical medications including topical corticosteroids, topical acyclovir and sometimes internal medications such as oral corticosteroids. Patients also tend to treat skin lesions with many over-the-counter medications such. . .

L11 ANSWER 12 OF 22 USPATFULL on STN

ACCESSION NUMBER: 96:55744 USPATFULL <<LOGINID::20080928>>

TITLE: Hyaluronic acid-urea pharmaceutical compositions and

uses

Gallina, Damian J., Erie, PA, United States INVENTOR(S):

PATENT ASSIGNEE(S): Patent Biopharmaceutics, Inc., Erie, PA, United States

(U.S. corporation)

NUMBER KIND DATE -----

PATENT INFORMATION: US 5529987 19960625
APPLICATION INFO.: US 1995-471331 19950602 (8)

RELATED APPLN. INFO.: Division of Ser. No. US 1993-101826, filed on 4 Aug

1993

Utility DOCUMENT TYPE: FILE SEGMENT: Granted

PRIMARY EXAMINER: Lilling, Herbert J. LEGAL REPRESENTATIVE: Cushman Darby & Cushman

NUMBER OF CLAIMS: EXEMPLARY CLAIM: 1

NUMBER OF DRAWINGS: 6 Drawing Figure(s); 6 Drawing Page(s)

CAS INDEXING IS AVAILABLE FOR THIS PATENT.

AB . . . erythema, edema, papules, vesicles, macules, pustules, scaling, cracking, crusting, and lesions. The invention further includes methods for the treatment of <u>psoriasis</u>, eczema, dermatitis, herpetic conditions, acne, skin ulcers, genital herpes lesions and anorectal disease, which includes applying to tissues in need. . .

. . . routinely in a dermatologic office setting. Normally, these conditions have been treated with various topical medications including topical corticosteroids, topical <u>acyclovir</u> and sometimes internal medications such as oral corticosteroids. Patients also tend to treat skin lesions with many over-the-counter medications such. . .

L11 ANSWER 13 OF 22 EMBASE COPYRIGHT (c) 2008 Elsevier B.V. All rights reserved on STN

ACCESSION NUMBER: 1996358293 EMBASE <<LOGINID::20080928>> TITLE: Herpes Zoster with unusual clinical course.

AUTHOR: Mazzotta, F., Dr. (correspondence); Troccoli, T.; Garofalo, L.; Bonifazi, E.

Clinica Dermatologica, Policlinico, University of Bari, CORPORATE SOURCE:

Piazza G. Cesare 11, 70124 Bari, Italy.

European Journal of Pediatric Dermatology, (1996) SOURCE:

Vol. 6, No. 1, pp. 25-30.

ISSN: 1122-7672 CODEN: EPDDE9

COUNTRY: Italy

DOCUMENT TYPE: Journal; Article

FILE SEGMENT: 013 Dermatology and Venereology

> 037 Drug Literature Index

004 Microbiology: Bacteriology, Mycology, Parasitology

and Virology

English LANGUAGE: SUMMARY LANGUAGE: English

ENTRY DATE: Entered STN: 9 Jan 1997

Last Updated on STN: 9 Jan 1997

European Journal of Pediatric Dermatology, (1996) Vol. 6, No. 1,

pp. 25-30.

ISSN: 1122-7672 CODEN: EPDDE9

. . . associated with moderate pain of the scapular-humeral joint. These symptoms and signs led to diagnose herpes zoster and to prescribe

Acyclovir per mouth. One month later, the patient came back

reporting a flare up of the lesions seven days after their. . . to the

diagnosis, due to the occurrence of some psoriatic scales. A

capillaroscopic study confirmed the diagnosis of eruptive zoster-like

psoriasis suggesting a Koebner phenomenon induced by herpes

(aciclovir) 59277-89-3; (salicylic acid) 63-36-5, 69-72-7 RN

L11 ANSWER 14 OF 22 USPATFULL on STN

95:7699 USPATFULL <<LOGINID::20080928>> ACCESSION NUMBER:

TITLE: Anti-inflammatory formulations for inflammatory

diseases

Kross, Robert D., Bellmore, NY, United States INVENTOR(S):

Siff, Elliott J., Westport, CT, United States

Alcide Corporation, Norwalk, CT, United States (U.S. PATENT ASSIGNEE(S):

corporation)

NUMBER KIND DATE ______

US 5384134 US 1993-115461 PATENT INFORMATION: 19950124

APPLICATION INFO.: 19930901 (8)

RELATED APPLN. INFO.: Continuation of Ser. No. US 1992-930088, filed on 14

Aug 1992, now abandoned which is a division of Ser. No. US 1990-543655, filed on 26 Jun 1990, now abandoned which is a division of Ser. No. US 1988-202758, filed on 3 Jun 1988, now patented, Pat. No. US 4956184 which is a continuation-in-part of Ser. No. US 1988-190798,

filed on 6 May 1988, now abandoned

DOCUMENT TYPE: Utility FILE SEGMENT: Granted

Cintins, Marianne M. PRIMARY EXAMINER:

Criares, T. J. ASSISTANT EXAMINER: LEGAL REPRESENTATIVE: Seed and Berry

NUMBER OF CLAIMS: EXEMPLARY CLAIM:

NUMBER OF DRAWINGS: 4 Drawing Figure(s); 4 Drawing Page(s)

LINE COUNT: 764

CAS INDEXING IS AVAILABLE FOR THIS PATENT.

AB

```
fungal infections, eczema, dandruff, acne, genital herpes lesions, and
       leg ulcers. There is further disclosed an antiviral lubricating
       composition that. . .
       . . . and secondary attacks become less frequent with time.
SUMM
       Treatments include drying agents to symptomatically lessen the
       discomfort of the lesion. Acyclovir, applied topically, tends
       to decrease pain of the primary lesions, but it has not proven very
       effective for decreasing vital shedding or lesion duration. Topical
       acyclovir has not been shown to be particularly effective for
       reducing or treating recurrent disease.
       Acyclovir is a purine nucleoside analog that is selectively
       cidal to the herpes simplex virus because only the thymidine kinase
       enzyme of herpes simplex virus can convert acyclovir to its
       monophosphate form while host cell thymidine kinase cannot. The
       monophosphate form is converted to an acyclovir triphosphate,
       which can interfere with vital DNA replication. Topical
       acyclovir is applied as a 5% ointment every three hours, or up
       to eight times daily, for at least seven days.. . . patient
       compliance problems for dosing in the genital areas throughout the day
       and throughout the night. A further problem of acyclovir has
       been the development resistant strains of herpes simplex, caused by a
      mutation of the thymidine kinase gene. Accordingly, no backup treatments
       are available for acyclovir-resistant herpes simplex
       infections. This problem exists with most antibiotic microbial
       treatments, but is generally not a problem non-antibiotic treatments.
DETD
       . . . as re-epithelialization of the original lesions). The results
       of the study were compared to a similar study conducted with topical
       acyclovir and placebo (Fiddian et al, J. Antimicrob. Chem.
       12:Suppl. B:67-77, 1983) and are presented together in Table 1 below:
       . . . Duration of
DETD
              Viral
                        Median
                                 Recurrence
       Symptoms
               Shedding Healing Rate Time (d) Time (d) %
       (d)
Example 1
                   1 * *
                             8 (1-17)
                                    19.4
(32)
 Acyclovir
                   3
                             7-8
                                    35
         5
                             10-13 55
Placebo 8
                   6-9
 *Twenty-one of twentyfour patients had a duration of symptoms of 5 or.
DETD twice daily dosing (compared with 5 times daily with some treatments
       such as Acyclovir)
L11 ANSWER 15 OF 22
                         MEDLINE on STN
                                  MEDLINE <<LOGINID::20080928>>
ACCESSION NUMBER: 1995332542
                    PubMed ID: 7541811
DOCUMENT NUMBER:
TITLE:
                    Primary Sjogren's syndrome and psoriasis vulgaris in a case
                    of OKT4 epitope deficiency.
                    Tanaka H; Mizutani H; Okada H; Shimizu M
                    Department of Dermatology, Mie University Faculty of
CORPORATE SOURCE:
                    Medicine, Tsu, Japan.
SOURCE:
                    The Journal of dermatology, (1995 Apr) Vol. 22,
                    No. 4, pp. 262-6.
                    Journal code: 7600545. ISSN: 0385-2407.
```

There is disclosed a method for treating dermatologic diseases caused by

microbial overgrowth or inflammation, such as psoriasis,

PUB. COUNTRY: Japan DOCUMENT TYPE: (CASE REPORTS) Journal; Article; (JOURNAL ARTICLE) LANGUAGE: English FILE SEGMENT: Priority Journals ENTRY MONTH: 199508 Entered STN: 28 Aug 1995 ENTRY DATE: Last Updated on STN: 29 Jan 1996 Entered Medline: 17 Aug 1995 The Journal of dermatology, (1995 Apr) Vol. 22, No. 4, pp. SO Journal code: 7600545. ISSN: 0385-2407. We report a 29-year-old female OKT4 epitope deficiency patient with primary Sjogren's syndrome and psoriasis vulgaris. Immunological investigations during the prolonged clinical course of her herpes zoster revealed that she has OKT4 epitope deficiency and primary Sjogren's syndrome. She had been treated for psoriasis vulgaris for 17 years without systemic immunosuppressive therapy. Flow cytometric study revealed that her OKT4 deficiency is heterogeneous and excluded interference with the OKT4 epitope by anti OKT4 autoantibodies. The rare coexistence of primary Sjogren's syndrome and psoriasis implicates an immune disturbance due to an unusual phenotype of CD4. Check Tags: Female CTAcyclovir: TU, therapeutic use Adult *Antigens, CD4: IM, immunology Biopsy *Epitopes: IM, immunology Flow Cytometry Herpes Zoster: CO, complications Herpes. . . RN 59277-89-3 (Acyclovir) L11 ANSWER 16 OF 22 MEDLINE on STN ACCESSION NUMBER: 1994267277 MEDLINE <<LOGINID::20080928>> PubMed ID: 8207272 DOCUMENT NUMBER: AIDS and the gateway of the body. TITLE: Bandyopadhyay P; Bhowal R N; Sikdar S N; Roy A K; Roy J G; AUTHOR: Bandyopadhyay D; Pal N C; Chatterjee B D CORPORATE SOURCE: Department of Dental Surgery, Medical College, Calcutta. SOURCE: Journal of the Indian Medical Association, (1994 Jan) Vol. 92, No. 1, pp. 17-9. Journal code: 7505608. ISSN: 0019-5847. Report No.: PIP-099042; POP-00233840. PUB. COUNTRY: India DOCUMENT TYPE: Journal; Article; (JOURNAL ARTICLE) LANGUAGE: English FILE SEGMENT: Priority Journals; Population; AIDS ENTRY MONTH: 199407 ENTRY DATE: Entered STN: 21 Jul 1994 Last Updated on STN: 1 Nov 2002 Entered Medline: 8 Jul 1994 SO Journal of the Indian Medical Association, (1994 Jan) Vol. 92, No. 1, pp. 17-9. Journal code: 7505608. ISSN: 0019-5847. Report No.: PIP-099042; POP-00233840. . . lesions in HIV-infected persons are larger and more numerous than

those in children. Various cutaneous or noncutaneous noninfective conditions (e.g., <u>psoriasis</u> and vasculitis) are also more common in HIV-infected persons. Possible agents to control candidiasis are

fluconazole and chlorhexidine oral rinse. Topical or systemic corticosteroids may control aphthous-like ulcers. The drug acyclovir may control herpes virus and other viral infections. If acyclovir is ineffective, desciclovir, ganciclovir, or foscarnet are possible alternatives. Papilloma virus lesions can be treated with cryosurgery, laser therapy, or.

L11 ANSWER 17 OF 22 EMBASE COPYRIGHT (c) 2008 Elsevier B.V. All rights reserved on STN

ACCESSION NUMBER: 1994014424 EMBASE <<LOGINID::20080928>>

TITLE: Liposomes: A promising future for dermatocosmetology and

clinical dermatology.

Raskovic, D. (correspondence); Piazza, P. AUTHOR:

CORPORATE SOURCE: Isto. Dermopatico dell'Immacolata, IRCCS, Via Monti di

Creta 104, 00167 Rome, Italy.

SOURCE: Journal of Liposome Research, (1993) Vol. 3, No.

3, pp. 737-751.

ISSN: 0898-2104 CODEN: JLREE7

COUNTRY: United States

Journal; Conference Article; (Conference paper) DOCUMENT TYPE:

FILE SEGMENT: 013 Dermatology and Venereology

> Clinical and Experimental Biochemistry 029

037 Drug Literature Index

LANGUAGE: English SUMMARY LANGUAGE: English

ENTRY DATE: Entered STN: 30 Jan 1994

Last Updated on STN: 30 Jan 1994

Journal of Liposome Research, (1993) Vol. 3, No. 3, pp. 737-751. ISSN: 0898-2104 CODEN: JLREE7

. . . capillary alterations. Liposomes have proven to be very useful AB for the therapy of certain dermatoses, such as atopic dermatoses or psoriasis. The advantage of incorporating a pharmaceutical substance (antibiotic, cortisone, immunomodulator, antimycotic, antiviral) can be observed in more effective and shorter therapy together with a decrease of side effects both local and linked to the systemic assimilation. Studies with acyclovir, interferon and topic steroids (triamcinolone and hydrocortisone) have been carried out experimentally. It is certain that a substance will have.

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ACCESSION NUMBER: 1993103284 EMBASE <<LOGINID::20080928>>

New drugs in pediatric dermatology. TITLE:

AUTHOR: Davis, A.; Krafchik, B.R. (correspondence)

CORPORATE SOURCE: 600 Sherbourne Street, Toronto, Ont. M4X 1W4, Canada.

Current Opinion in Pediatrics, (1993) Vol. 5, No. SOURCE:

2, pp. 212-215.

ISSN: 1040-8703 CODEN: COPEE9

COUNTRY: United States

DOCUMENT TYPE: Journal; General Review; (Review) FILE SEGMENT: 013 Dermatology and Venereology

> 037 Drug Literature Index 038 Adverse Reactions Titles

007 Pediatrics and Pediatric Surgery

LANGUAGE: English SUMMARY LANGUAGE: English

ENTRY DATE: Entered STN: 16 May 1993

Last Updated on STN: 16 May 1993

SO Current Opinion in Pediatrics, (1993) Vol. 5, No. 2, pp. 212-215.

```
ISSN: 1040-8703 CODEN: COPEE9
     . . . infectious and inflammatory skin diseases. Five of these drugs,
AB
     calcipotriol, EMLA (eutectic mixture of local anesthetics),
     interferon-\alpha 2a, cyclosporine, and <u>acyclovir</u>, are
     reviewed. Calcipotriol, a vitamin D analogue, has been shown to be useful
     but not curative in chronic stable plaque psoriasis in adults.
     Its use in children is being studied. EMLA is proving to be useful in the
     prevention of pain. . . of vascular lesions. Cyclosporine is a
     powerful immunomodulating agent. It has been used in the treatment of
     atopic dermatitis and psoriasis. Adverse effects limit its
     widespread use as a systemic agent. Topical cyclosporine has limited
     efficacy due to its poor penetration. Acyclovir has
     revolutionized the treatment of herpes simplex virus infections,
     particularly in reducing the morbidity and mortality of neonatal herpes.
     Whether. .
RN
     (aciclovir) 59277-89-3; (alpha2a interferon) 76543-88-9;
     (calcipotrio1) 112828-00-9, 112965-21-6; (cyclosporin) 79217-60-0; (EMLA)
     101362-25-8
L11 ANSWER 19 OF 22 EMBASE COPYRIGHT (c) 2008 Elsevier B.V. All rights
     reserved on STN
                    1992176613 EMBASE <<LOGINID::20080928>>
ACCESSION NUMBER:
                     Results of treatment of genital herpes with
TITLE:
                     acyclovir produced by Polfa Pharmaceutical Factory
                     in Stargard.
AUTHOR:
                     Wojas, K. (correspondence); Szmacinska, E.; Pieprzny, H.;
                     Pilat, I.; Antonowicz, A.
CORPORATE SOURCE:
                     Wojewodzki Oddział Dermatologiczny, ul. Staszica 4A, 37-450
                     Stalowa Wola, Poland.
                     Przeglad Dermatologiczny, (1992) Vol. 79, No.
SOURCE:
                     1-2, pp. 65-69.
                     ISSN: 0033-2526 CODEN: PRDEA7
COUNTRY:
                     Poland
DOCUMENT TYPE:
                     Journal; Article
FILE SEGMENT:
                     013
                            Dermatology and Venereology
                     037
                             Drug Literature Index
LANGUAGE:
                     Polish
SUMMARY LANGUAGE:
                    English; Polish
ENTRY DATE:
                    Entered STN: 5 Jul 1992
                    Last Updated on STN: 5 Jul 1992
     Results of treatment of genital herpes with acyclovir produced
TΙ
     by Polfa Pharmaceutical Factory in Stargard.
     Przeglad Dermatologiczny, (1992) Vol. 79, No. 1-2, pp. 65-69.
     ISSN: 0033-2526 CODEN: PRDEA7
AB
     . . . patients underwent PUVA treatment and other 30 patients SUP
     treatment, with external assist therapy. As for the clinical forms of
     psoriasis patients with common psoriasis were the most
     numerous group. During PUVA therapy patients were irradiated 3-4 times a
     week with average number of exposures. . . early complications in both methods were observed in 4 patients (6.7\%). Both methods are valuable
     completion of classical treatment of \underline{\text{psoriasis}} . The differences between the results of the treatments were very small.
L11 ANSWER 20 OF 22 DISSABS COPYRIGHT (C) 2008 ProQuest Information and
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                     91:20834 DISSABS
                                        Order Number: AAR9201964
ACCESSION NUMBER:
                     TRANSDERMAL TRANSPORT AND INTRADERMAL DRUG TARGETING USING
TITLE:
                     NOVEL CHEMICAL DELIVERY SYSTEMS
AUTHOR:
                     CHIKHALE, PRASHANT JAYANT [PH.D.]; BODOR, NICHOLAS S.
```

[advisor]

CORPORATE SOURCE: UNIVERSITY OF FLORIDA (0070)

SOURCE: Dissertation Abstracts International, (1991) Vol.

52, No. 8B, p. 4209. Order No.: AAR9201964. 161 pages.

DOCUMENT TYPE: Dissertation

FILE SEGMENT: DAI LANGUAGE: English

Entered STN: 19921118 ENTRY DATE:

Last Updated on STN: 19921118

Dissertation Abstracts International, (1991) Vol. 52, No. 8B, p.

4209. Order No.: AAR9201964. 161 pages.

This dissertation examines the feasibility of targeting and AB localizing important antiviral agents (like acyclovir) and anticancer agents (like 5-fluorouracil) specifically to the skin using novel redox-based chemical targeting systems. Such approaches should lead to improvement in the effectiveness of topically administered acyclovir in treating recurrent mucocutaneous herpes simplex virus infection of type I. Similarly, the basal cell skin carcinomas or psoriasis can be effectively treated if 5-fluorouracil could be targeted to the intra-dermal region.

Chemical Delivery Systems (CDS) for acyclovir based on oxidation (the 1,4-dihydrotrigonelline moiety containing ester; A-CDS) or reduction (the lipoic acid ester; A-LipS\$\sb2\$) in the skin were utilized to enhance the skin-partitioning ability of acyclovir and use the enzymatic activity of the skin to create metabolic chemical precursors as reservoirs for the release of acyclovir in the skin.

Thus, the dermal delivery of acyclovir was improved by 9-fold (p < 0.025) using A-CDS, and by 37-fold (p < 0.001) using A-LipS\$\sb2\$, at 6 hours relative to underivatized acyclovir, when administered to the hairless-mouse skin, in vitro. The lipolyl ester of 5-fluorouracil (5-FU-LipS\$\sb2\$) also managed to deliver greater 5-fluorouracil. . .

L11 ANSWER 21 OF 22 USPATFULL on STN

ACCESSION NUMBER: 90:71577 USPATFULL <<LOGINID::20080928>> TITLE: Topical treatment of genital herpes lesions
INVENTOR(S): Kross, Robert D., Bellmore, NY, United States
PATENT ASSIGNEE(S): Alcide Corporation, Norwalk, CT, United States (U.S.

corporation)

NUMBER KIND DATE ______

PATENT INFORMATION: US 4956184 19900911 APPLICATION INFO.: US 1988-202758 19880603 (7)

RELATED APPLN. INFO.: Continuation-in-part of Ser. No. US 1988-190798, filed

on 6 May 1988, now abandoned Utility

DOCUMENT TYPE:

FILE SEGMENT: Granted
PRIMARY EXAMINER: Friedman, Stanley J.
LEGAL REPRESENTATIVE: Seed and Berry
NUMBER OF CLAIMS:

NUMBER OF CLAIMS: EXEMPLARY CLAIM: 1

NUMBER OF DRAWINGS: 4 Drawing Figure(s); 2 Drawing Page(s) LINE COUNT: 715

CAS INDEXING IS AVAILABLE FOR THIS PATENT.

There is disclosed a method for treating dermatologic diseases caused by microbial overgrowth or inflammation, such as psoriasis, fungal infections, eczema, dandruff, acne, genital herpes lesions, and leg ulcers. There is further disclosed an antiviral lubricating composition that. . .

SHMM

```
. . . and secondary attacks become less frequent with time.
       Treatments include drying agents to symptomatically lessen the
       discomfort of the lesion. <a href="Acyclovir">Acyclovir</a>, applied topically, tends
       to decrease pain of the primary lesions, but it has not proven very
       effective for decreasing viral shedding or lesion duration. Topical
       acyclovir has not been shown to be particularly effective for
       reducing or treating recurrent disease.
SUMM
       Acyclovir is a purine nucleoside analog that is selectively
       cidal to the herpes simplex virus because only the thymidine kinase
       enzyme of herpes simplex virus can convert acyclovir to its
       monophosphate form while host cell thymidine kinase cannot. The
       monophosphate form is converted to an acyclovir triphosphate,
       which can interfere with viral DNA replication. Topical acycovir is
       applied as a 5% ointment every three hours, or. . . patient
       compliance problems for dosing in the genital areas throughout the day
       and throughout the night. A further problem of acyclovir has
       been the development of resistant strains of herpes simplex, caused by a
       mutation of the thymidine kinase gene. Accordingly, no backup treatments
       are available for acyclovir-resistant herpes simplex
       infections. This problem exists with most antibiotic antimicrobial
       treatments, but is generally not a problem for non-antibiotic
       treatments.
       . . . as re-epithelialization of the original lesions). The results
DETD
       of the study were compared to a similar study conducted with topical
       acyclovir and placebo (Fiddian et al., J. Antimicrob. Chem.
       12:Suppl. B:67-77, 1983) and are presented together in Table 1 below:
       . . . Duration of
DETD
              Viral
                        Median
                                Recurrence
       Symptoms
               Shedding Healing Rate
       (d)
               Time (d) Time (d) %
Example 1
          3*
                   1 * *
                             8 (1-17)
                                    19.4
(32)
  Acyclovir
                                    35
                   3
                             7-8
Placebo 8
                   6-9
                             10-13 55
 *Twenty-one of twentyfour patients had a duration of symptoms of 5 or.
      twice daily dosing (compared with 5 times daily with some treatments
       such as Acyclovir)
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     reserved on STN
ACCESSION NUMBER:
                   1990140613 EMBASE
                                        <<LOGINID::20080928>>
TITLE:
                    Common cutaneous disorders in athletes.
AUTHOR:
                    Conklin, R.J.
CORPORATE SOURCE:
                    Department of Dermatology, University of British Columbia,
                    Vancouver, BC, Canada.
                    Sports Medicine, (1990) Vol. 9, No. 2, pp.
SOURCE:
                    100-119.
                    ISSN: 0112-1642 CODEN: SPMEE7
```

COUNTRY:

LANGUAGE:

DOCUMENT TYPE: FILE SEGMENT:

SUMMARY LANGUAGE:

New Zealand

035

English

English

Journal; General Review; (Review)

Dermatology and Venereology

Occupational Health and Industrial Medicine

ENTRY DATE: Entered STN: 13 Dec 1991

Last Updated on STN: 13 Dec 1991

Sports Medicine, (1990) Vol. 9, No. 2, pp. 100-119. ISSN: 0112-1642 CODEN: SPMEE7 SO

AΒ . . . be treated with any drying agents (e.g. alcohol) as they are as effective as more expensive topical agents such as acyclovir. Molluscum contagiosum may be spread by close contact or water contact and is treated by superficial incision, cryotherapy or standard. . . normal foot function and minimal surgical procedures. Paronychia is treated best by wedge resection. Sweat and friction may aggravate pre-existing psoriasis, acne, atopic dermatitis and allergic contact dermatitis. Allergic contact dermatitis may be caused by dyes, rubber chemicals or glues associated. .

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